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REMARKS

The Notice of Non-Compliant Amendment states that the claims filed on April 16, 2004 do not comply with the requirements of 37 C.F.R. § 1.121(c) because claims 28 and 29 are listed as “previously presented” but in fact are “currently amended.” In response, Applicants are submitting an Amendment which is substantially similar to the Amendment filed on April 16, 2004, but which has been amended to correct this inadvertent error.

After entry of this Amendment, claims 14, 16-20, 23-29, 31-32 and 35-37 will be pending and under consideration in this application.

Claims 14, 16, 19-20, 23-29 and 31-32 have been amended. Claims 35-41 have been added. Support for the claim amendments and for the new claims can be found throughout the specification. (The specific support for the claim amendments and new claims will be discussed below.) Applicants respectfully submit that none of the claim amendments add new matter to the application.

Claims 1, 3-4, 7-9 and 12-13 have been cancelled without prejudice. Applicants reserve the right to file an application which claims priority to the instant application and contains the subject matter of the cancelled claims.

Indefiniteness Rejections under 35 U.S.C. §112, ¶2

The Examiner has rejected claims 1, 3, 4, 7-9, 12-14, 16-20, 23-29, 31 and 32 under 35 U.S.C. §112, ¶2, as being indefinite due to their recitation of M64347_at. The Examiner states that M64347_at is a GenBank Accession No. and therefore an object which is variable.

Claims 1, 3-4, 7-9 12 and 13 have been cancelled and, therefore, the rejection with respect to these claims has been obviated.

Applicants respectfully traverse the rejection with respect to claims 14, 16-20, 23-29, 31 and 32, which have been amended to refer to an informative gene comprising certain nucleotides of GenBank Accession No. M64347. Support for this amendment can be found at in the specification which discloses that the Affymetrix “HuGeneFL array” was used, and that M64347_at marker was upregulated. See Amendments to the Specification submitted in July 30, 2003, under the heading “Microarray hybridization” and Table 1. As indicated in the Affymetrix

website (Exhibit A), M64347_ at refers to a set of probes which detect nucleotides 3336-3720 of GenBank Accession No. M64347.

Applicants submit that reference to a GenBank Accession No. does not render the claims indefinite. A person of ordinary skill in the art would be able to determine the nucleotide sequence of a gene by reference to its GenBank Accession No. Further, while a GenBank Accession No. can be revised, a person of ordinary skill in the art would be able to identify any revisions made to a sequence over time. Attached as Exhibit B is print-out from the National Center for Biotechnology Information ("NCBI") website which summarizes their policy with respect to sequences revisions (*see* page 1 of the Exhibit), and states that a person reviewing the records for a particular GenBank Accession No. would be able to determine whether a particular sequence has been revised and would be able to access previous versions of the sequence (*see* page 3 of the Exhibit).

Furthermore, a search of the United States Patent and Trademarks Office ("USPTO") granted patents database reveals that the USPTO has granted patents containing claims which recite sequences by references to their GenBank Accession Nos.¹ Thus, Applicants respectfully request that the Examiner withdraw this rejection.

The Examiner has also rejected claims 12-13, 24-25, 28 and 29 for referring to Table 1 and Tables 2-6. Further, the Examiner states that the reference to the genes in Tables 1-6 by reference to their GenBank Accession Nos. renders the claims indefinite.

Claims 12 and 13 have been cancelled and, therefore, the rejection with respect to these claims has been obviated.

Claims 24-25, 28 and 29 have been amended to replace the reference to the Tables with reference to the GenBank Accession Nos. for the genes disclosed in the Table. Support for this amendment can be found at Tables 1 and 6. Table 1 lists the genes by reference to the probes used to detect the genes disclosed in Table 1, which correlate to the GenBank Accession Nos. for the genes disclosed in the Table 1. Exhibit C provides information obtained from the Affymetrix website which shows such correlation. Note that Exhibit C does not provide the information for all of the probes disclosed in Table 1 since the purpose of the exhibit is to

¹ The query used was: "GenBank Accession Number" or "GenBank Accession No." in claims. This search resulted in 18 hits. Among the relevant hits were: U.S. Patent Nos. 6,667,065, 6,627,193, 6,468,773, and others.

demonstrate that the reference to the probes in Table 1 correlates with the GenBank Accession Nos. of the genes disclosed in Table 1.

Applicants traverse the Examiner's statement that references to GenBank Accession Nos. render the claims indefinite for the reasons discussed above. Accordingly, Applicants respectfully submit that the scope of claims 24-25, 28 and 29 is definite, and request the Examiner to withdraw this rejection.

Applicants note that claims 40 and 41 have been added. Support for the claims can be found throughout the specification, particularly at Tables 1 and 6. Applicants respectfully submit that claims 40 and 41 are definite.

The Examiner has rejected claim 1 as indefinite "for failing to [show the] how the expression profile is correlated with a specific brain tumor type." Further, the Examiner states that "the metes and bounds of two or more informative genes beyond the M64347" is unclear. Claim 1 has been cancelled and, therefore, this rejection has been obviated.

The Examiner has rejected claims 8, 9, 19 and 20 as indefinite because they "lack active method steps, as the recitation of 'utilizing' does not constitute a specific method step."

Claims 8 and 9 have been cancelled and, therefore, the rejection with respect to these claims has been obviated. Applicants respectfully traverse this rejection with respect to claims 19 and 20. Applicants submit that the recitation of "utilizing" in claims 19 and 20 is not intended to be a further method step. Instead the recitation of utilizing provides a further definition of how to determine a gene expression profile. Applicants have amended the claims to improve their form and clarify this point. Accordingly, Applicants respectfully request that the Examiner withdraw this rejection.

The Examiner has rejected claim 14 as indefinite because the use of the term "the sample" lacks antecedent basis. Claim 14 as amended refers to "the brain tumor" rather than "the sample." This amendment renders the rejection moot.

The Examiner has rejected claim 23 as indefinite for its recitation of "survival after treatment" as the predicted treatment outcome. Applicants have amended claim 23 to require the predicted treatment outcome to be a good prognosis of survival after treatment or treatment failure. Support for amended claim 23 can be found at page 9, lines 23-26 of the Substitute Specification. This amendment renders the rejection moot.

The Examiner has rejected claims 26 and 27 as indefinite because of their recitation of: “informative genes”, “magnitude”, “class distinction”, “winning”, and “summing the votes”, “the sample to be tested”, “first class” and “second class”. (See Office Action, pages 3-4.) More specifically, with respect to claim 26, the Examiner states that “it is unclear how the ‘magnitude’ of the vote is to be determined because ‘depending on the expression level of the gene’ does not accurately define the mathematical relationship between the gene expression and the magnitude of the vote” (Office Action, page 3).

In response to the Examiner’s rejections of claims 26 and 27, Applicants traverse in part and amend in part. Applicants have amended these claims to improve their form and to more clearly define the claimed invention. Applicants respectfully submit that the amended claims would be considered definite by a person having ordinary skill in the art.

The claims as amended require calculating the weighted vote of each informative gene. According to the specification, “informative genes” refers to “genes whose expression correlates with a particular phenotype.” (See Substitute Specification, page 8, lines 23-25 and page 9, lines 13-16.) Thus, with respect to claims 26 and 27, an informative gene is one which correlates with treatment outcome.

Further, the claims have been amended to clarify “class distinction”, “first class” and “second class”. Applicants respectfully submit that based on the information disclosed in the specification and the knowledge in the art, a person of skill in the art at the time the application was filed would be able to calculate the weighted vote for an informative gene, and to sum up the votes to determine a winning class as required by claims 21 and 22. The weighted voting algorithm was well known in the art at the time the application was filed as evidenced by the fact that the specification cites to three references which use this method. See Substitute Specification, page 32, lines 27-28 and Amendments to the Specification submitted in July 30, 2003, under the heading “Weighted Voting”, citing to: U.S. Application No. 09/544,627 (now issued as U.S. Patent No. 6,647,341), Golub 1999, and Slonim 2000.

Finally, the Examiner states that claims 26 and 27 are vague and indefinite because it is unclear if the level of gene expression used in the computation is a normalized or non-normalized level. Applicants respectfully traverse. Applicants submit that it is irrelevant to the claimed methods, and that a person of skill in the art would know whether the gene expression

level should be normalized or non-normalized in a particular instance. Accordingly, Applicants respectfully request that the Examiner withdraw this rejection.

Enablement Rejections under 35 U.S.C. §112, ¶1

The Examiner has rejected claims 1, 3, 4, 7-9, 12-14, 16-20 and 23-29 for lack of enablement.

Applicants have cancelled claims 1, 3, 4, 7-9, 12 and 13, directed to methods of classifying a brain tumor. Therefore, the Examiner's rejections with respect to these claims have been obviated.

Applicants traverse the Examiner's enablement rejection with respect to claims 14, 16-20 and 23-29, directed to methods of predicting the efficacy of treating a brain tumor, methods for predicting a treatment outcome of a patient with a brain tumor, methods for evaluating drug candidates for their effectiveness in treating a brain tumor or methods for monitoring the efficacy of a brain tumor.

According to the Examiner, "[t]here are no teachings in the specification to correlate a value which is several standard deviations from the mean with a method of predicting the efficacy of a brain tumor. More specifically, the Examiner states that: (1) the specification does not teach whether the expression of M64347 as shown in Figure 3C was obtained from the brain tumor before treatment or after treatment, (2) it is unclear if the lowered expression of M64347 is indicative or predictive of treatment failure or a treatment success as the title of Figure 3C is "Markers of Treatment Failure" but the heading of Table 1 is "Markers Downregulated with Low Risk" and (3) the specification does not define how the C1 or C0 groups were differentiated and does not teach what constitutes a treatment failure or success in terms of disease free survival or length of survival.

As mentioned above, Applicants traverse. First, the specification teaches that expression of M64347 as shown in Figure 3C was obtained from the brain tumor before treatment. *See* Substitute Specification, page 40, line 28 to page 41, line 2.

Second, the specification (Table 1, Table 6 and Figure 3C) shows that the upregulation of M64347 is correlated with a "high risk class" of individuals (*e.g.*, a class of individuals with

poor prognosis for survival after treatment). *See* Substitute Specification, page 9, lines 13-16 stating that “a sample can be classified as belonging to a high risk class (e.g., a class with poor prognosis for survival after treatment) or a low risk class (e.g., a class with good prognosis for survival after treatment).” Thus, the heading of Table 1 – “Markers Upregulated in High Risk, Downregulated in low Risk” – is not inconsistent with the heading of Figure 3C – “Markers of Treatment Failure” – as it appears to be suggested by the Examiner.

Third, the specification describes that in Figure 3 C0 and C1 correspond to two unsupervised SOM-derived clusters, and that Class C1 tumors are notable for their high ribosomal content. *See* Substitute Specification, page 7, lines 2-4. The specification further states that the C0 and C1 groups were not correlated with patient survival. *See* Substitute Specification, page 41, lines 13-17.

Finally, the specification teaches what constitutes treatment failure or success in terms of disease free survival or length of survival. The specification states that they differentiated “patients who are alive following treatment (‘survivors’) compared to those [patients] who succumbed to their disease (‘failures’; minimum follow-up 24 months for surviving patients; overall median 41.5 months).” *See* Substitute Specification page 41, lines 17-21.

The Examiner also states that “[t]here is no guidance for a specific polynucleotide probe and hybridization conditions to be used in the determination of an expression profile for . . . the method of predicting the efficacy of treatment.” The Examiner noted there are different isoforms of FGFR3, the gene encoded by M64347, and stated that a probe to this gene could hybridize to any number of the polymorphic gene products or alleles. The Examiner concluded that the “specification provides no teachings as to the exact nature of the probe used for the expression profile, thus it cannot be construed from the specification which polymorphic variants, splice variants or alleles are integral to the claimed invention.”

Applicants note that not all of the claims recite the use of a probe and/or require the use of hybridization condition to determine an expression profile. In any case, Applicants respectfully traverse the Examiner’s rejection to the extent that certain claims require the use of a specific polynucleotide probe and hybridization conditions.

Contrary to the Examiner’s assertion, the specification teaches the exact nature of the probes used to determine the expression profiles for the classification of a brain tumor, or the

method of predicting the efficacy of treatment. The specification states that they used Affymetrix's HuGeneFL array. (See Amendments to the Specification submitted in July 30, 2003, under the heading "Microarray hybridization.") Based on this disclosure, a person of skill in the art would have been able to identify the probes present in the array and used in the specification to determine the expression profiles for the method of predicting the efficacy of treating a brain tumor.

More specifically, based on the disclosure, a person skilled in the art would have been able to determine the specific probe used to determine the expression profile of M64347, and the other informative genes disclosed in the specification. The claims have been amended to recite an informative gene comprising nucleotides 3336-3720 of GenBank Accession No. M64347, which are the nucleotide sequences in Affymetrix's HuGeneFL array. See Exhibit A (obtained from Affymetrix's website). Thus, Applicants submit that the claims, as amended, are enabled with respect to the probes which can be used to practice the methods of the claimed invention.

Applicants respectfully submit that, in view of the specification, which teaches that the expression profile of an informative gene which hybridizes to nucleotides 3336 to 3720 of GenBank Accession No. M64347 correlates with efficacy of treating a brain tumor, the fact that there are different allelic variants or isoforms of the gene encoded by GenBank Accession No. M64347 is irrelevant.

Applicants respectfully submit that a person of ordinary skill in the art would know what hybridization conditions should be employed to determine the expression profile of the informative genes of the claims. Moreover, the specification teaches the hybridization conditions used in the experiments disclosed in the specification. (See Amendments to the Specification submitted in July 30, 2003, under the heading "Microarray hybridization.") Thus, Applicants submit that the claims are enabled with respect to the hybridization conditions useful in practicing the methods of the claimed invention.

With respect to claims 26-29, the Examiner states that "the specification does not define the parameters needed to calculate weighted vote for M64347." Applicants respectfully traverse. As discussed above, based on the information disclosed in the specification, a person of skill in the art would know how to determine weighted vote as recited in the claims without undue experimentation as evidenced by the fact that the specification refers to a patent

application (U.S. Application No. 09/544,627, now issued as U.S. Patent No. 6,647,341) and two papers (Golub 1999 and Slonim 2000) that disclose the use of this weighted voting algorithm before the instant application was filed. (See Substitute Specification, page 32, lines 27-28 and Amendments to the Specification submitted in July 30, 2003, under the heading "Weighted Voting.")

The Examiner states that "Applicants arguments regarding the teachings of Golub et al. for methods of determining class and subclass as set forth in U.S. application No. 09/544,627 are unpersuasive" because the instant application could issue before the referenced application. (See Office Action, page 7.) Applicants note that the referenced application has now issued as U.S. Patent No. 6,647,341. Applicants have amended the application accordingly.

In view of the arguments presented above, Applicants respectfully request that the Examiner withdraw the enablement rejections.

Obviousness Rejections

The Examiner has rejected claims 31 and 32 under 35 U.S.C. §103(a) as being unpatentable over U.S. Patent No. 6,500,938 ("*Au-Young*") in view of Abbass et al., 1997, *J. Clin. Endocrinol. Metab.*, 82:1160-1166 ("*Abbas*"), or over U.S. Patent No. 6,218,122 ("*Friend*") in view of *Abbas*.

The Examiner alleges that *Au-Young* teaches methods of monitoring the progression of a disease or the efficacy of a treatment comprising detecting an expression profile by means of a micro array. The Examiner alleges that *Friend* teaches methods for detecting changes in a biological state of a subject which are correlated to one or more disease states and methods for monitoring the efficacies of a therapy comprising the determination of an expression profile from said cells in a patient. However, as admitted by the Examiner, neither *Au-Young* nor *Friend* teaches the expression profile of M64347 or the FGFR3 encoded thereby.

The Examiner alleges that *Abbas* teaches "that the expression of the mRNA encoding the secreted form of FGFR3, which would be expressed from the M64347_at gene, is correlated with pituitary adenomas." Further, as admitted by the Examiner, *Abbas* does not teach a correlation between the expression profile of FGFR3 and tumor type, size or aggressiveness. (See 12/31/03 Office Action, page 5.)

It is respectfully pointed out to the Examiner that a proper rejection based on 35 U.S.C. §103 that relies on a combination of prior art references requires a teaching, suggestion, or motivation to combine the teachings of the references; a reasonable expectation of success founded in the cited art of producing the claimed invention; and that such proper combination teaches or suggests all elements of the claimed invention. Applicants respectfully traverse the obviousness rejections for failing to meet all of these requirements for the reasons provided below.

Claims 31 and 32, as amended, recite methods for evaluating drug candidates for their effectiveness in treating brain tumors or methods for monitoring the efficacy of a brain tumor treatment, wherein the brain tumor is selected from the group consisting of melanoblastomas, glioblastomas, rhabdoid tumors, primitive neuroectodermal tumors, and pineoblastomas. Support for this amendment, and for newly added claims 35-39, can be found throughout the specification. *See, e.g.*, Substitute Specification, page 3, lines 1-2.

Applicants respectfully submit that there is no motivation to combine the references cited by the Examiner to reach the invention of amended claims 31 and 32. The Examiner states that “one of ordinary skill in the art would have been motivated to [combine the references] with a reasonable expectation of success by the teachings of Abbas et al. on the unique expression of the secreted form of FGFR3 mRNA in pituitary adenomas versus the lack of expression of the secreted form of this receptor in normal pituitary.” However, none of the references teach the correlation between the gene expression profile of M64347 and effectiveness in treating a brain tumor selected from the group consisting of melanoblastomas, glioblastomas, rhabdoid tumors, primitive neuroectodermal tumors, and pineoblastomas, or monitoring the efficacy of a treatment for any of the mentioned brain tumors. Therefore, there would be no motivation to combine the references as argued by the Examiner.

Further, even if the references were to be combined as suggested by the Examiner, the combination of references would not teach or suggest the inventions of claims 31 and 32. Rather, the combination of the references would at best teach the use of M64347 to evaluate drug candidates for their effectiveness in treating a pituitary adenoma, or to monitor the efficacy of a pituitary adenoma treatment. Nothing in the cited references suggests or discloses the use of M64347 to evaluate drug or monitor the efficacy of a drug to treat the brain tumors of the claims. Moreover, even if the references were combined, there would be no reasonable

expectation of success. Accordingly, Applicants respectfully request that the Examiner withdraw the obviousness rejections.

Conclusion

In view of the above, each of the presently pending claims in this application is believed to be in immediate condition for allowance. Accordingly, the Examiner is respectfully requested to pass this application to issue.

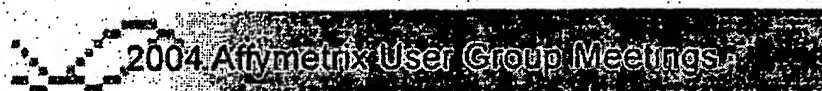
Applicant believes no fee is due with this response. However, if a fee is due, please charge our Deposit Account No. 18-1945, under Order No. WIBL-P01-561 from which the undersigned is authorized to draw.

Dated: July 15, 2004

Respectfully submitted,

By 
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Expression
→ Quick Query
→ Standard Query
→ Batch Query
→ BLAST
→ Probe Match
→ UCSC Query

Genotyping
→ Quick Query
→ Standard Query
→ Batch Query
→ UCSC Query
→ SNP Finder

≡ QUERY HISTORY
Annotation Views
→ Expression
→ Genotyping
→ BLAST Status

→ New Folder
→ Expression
Queries
→ (1) All Descriptions
(m64347)
→ all probe sets
(7129)

Full Record

Details for HUGENEFL:M64347_AT

Full Screen

NetAffx Links [Cluster Members](#)
[Consensus/Exemplar](#)

GeneChip Array Information

Probe Set ID M64347_at
GeneChip Array HumanGeneFL Array
Organism Common Name Human

Probe Design Information

Transcript ID M64347
Sequence Type Exemplar sequence
Representative Public ID M64347 [NCBI](#)
Target Description M64347, class A, 20 probes, 20 in M64347 3336-3720, Human novel growth factor receptor mRNA, 3' cds

Genomic Alignment of Target Sequence

Assembly April 2003 (NCBI 33)
Alignment(s) Position % Identity Cytoband
chr4: 1771773-1772182 (+) [UCSC](#) 93 p16:3

Representative Transcript	UniGene Description	Position
NM_000142 NCBI	fibroblast growth factor receptor 3 (achondroplasia, thanatophoric dwarfism)	chr4:1757261-1772237 (+) UCSC
NM_022965 NCBI	fibroblast growth factor receptor 3 (achondroplasia, thanatophoric dwarfism)	chr4:1757261-1772237 (+) UCSC

Public Domain and Genome References

Gene Title fibroblast growth factor receptor 3 (achondroplasia, thanatophoric dwarfism)
Gene Symbol FGFR3 [HGNC](#)
Chromosomal Location 4p16.3
UniGene ID Hs.1420 [NCBI](#) (FULL LENGTH)
Ensembl ENSG00000068078 [Ensembl](#)
LocusLink 2261 [NCBI](#)
P22607 [EMBL-EBI](#)

SwissProt	Q96T34 EMBL-EBI Q96T35 EMBL-EBI Q96T36 EMBL-EBI Q9NRB6 EMBL-EBI
EC	2.7.1.112
OMIM	134934 NCBI
RefSeq Protein ID	NP_000133 NCBI NP_075254 NCBI
RefSeq	RefSeq Transcript ID RefSeq Title NM_000142 NCBI fibroblast growth factor receptor 3 isoform 1 precursor NM_022965 NCBI fibroblast growth factor receptor 3 isoform 2 precursor

Functional Annotations

	ID	Title	Organism	Type
	DROSGENOME1:143549 AT	breathless	Drosophila	Putative Ortholog
	RAE230A:1369373 AT	fibroblast growth factor receptor 3	Rat	Putative Ortholog
	RAE230B:1384056 AT	fibroblast growth factor receptor 3	Rat	Putative Ortholog
	RAE230B:1384829 AT	fibroblast growth factor receptor 3	Rat	Putative Ortholog
	RG-U34B:RC AA899336 AT	fibroblast growth factor receptor 3	Rat	Putative Ortholog
	RG-U34C:RC AI136304 AT	fibroblast growth factor receptor 3	Rat	Putative Ortholog
	RG-U34C:RC AI145424 AT	fibroblast growth factor receptor 3	Rat	Putative Ortholog
Ortholog	MG-U74AV2:160919 R AT	fibroblast growth factor receptor 3	Mouse	Curated Ortholog
	MG-U74AV2:162253 I AT	fibroblast growth factor receptor 3	Mouse	Curated Ortholog
	MOE430A:1421841 AT	fibroblast growth factor receptor 3	Mouse	Curated Ortholog
	MOE430A:1425796 A AT	fibroblast growth factor receptor 3	Mouse	Curated Ortholog
	MU11KSUBA:M81342 S AT	fibroblast growth factor receptor 3	Mouse	Curated Ortholog
	MOUSE430 2:1421841 AT	fibroblast growth factor receptor 3	Mouse	Curated Ortholog
	MOUSE430 2:1425796 A AT	fibroblast growth factor receptor 3	Mouse	Curated Ortholog
	MOUSE430A 2:1421841 AT	fibroblast growth factor receptor 3	Mouse	Curated Ortholog
	MOUSE430A 2:1425796 A AT	fibroblast growth factor receptor 3	Mouse	Curated Ortholog

GO Biological Process (view graph)

ID	Description	Evidence	Links
165	MAPKKK cascade	experimental evidence	QuickGO AmiGO
1501	skeletal development	predicted/computed	QuickGO AmiGO
7048	oncogenesis	experimental evidence	QuickGO AmiGO
7259	JAK-STAT cascade	experimental	QuickGO

	8543 FGF receptor signaling pathway	evidence experimental evidence	AmiGO QuickGO AmiGO
	GO Cellular Component (view graph)		
	ID	Description	Evidence
Gene Ontology	5887 integral to plasma membrane	experimental evidence	QuickGO AmiGO
	GO Molecular Function (view graph)		
	ID	Description	Evidence
	5007 fibroblast growth factor receptor activity	experimental evidence	QuickGO AmiGO
Protein Similarities	Method	ID	Description
	blast	13112048	fibroblast growth factor receptor 3 isoform 2 precursor; hydroxyaryl-protein kinase; tyrosine kinase JTK4 [Homo sapiens]
	blast	13186255	fibroblast growth factor receptor 2 isoform 3 precursor; keratinocyte growth factor receptor; K-sam protein; protein tyrosine kinase, receptor like 14; FGF receptor; bacteria-expressed kinase; fibroblast growth factor receptor BEK; tyrosylprotein kinase; hydroxyaryl-protein kinase [Homo sapiens]
	blast	4503711	fibroblast growth factor receptor 3 isoform 1 precursor; hydroxyaryl-protein kinase; tyrosine kinase JTK4 [Homo sapiens]
Protein Families	blast	20452380	0.0
	Method	ID	Description
	Hanks	FGFR-3	FGR3_HUMAN (FGFR-3) KINASES:5.6.3 PTK Group B membrane spanning protein tyrosine kinases.PTK XV Fibroblast growth factor receptor family .FGFR-3
	ec	ZA70_HUMAN	ZA70_HUMAN EC:2.7.1.112:TYROSINE-PROTEIN KINASE ZAP-70 (EC 2.7.1.112) (70 KDA ZETA-ASSOCIATED PROTEIN) (SYK-RELATED TYROSINE KINASE).
	Hanks	FGFR-3	FGR3_HUMAN (FGFR-3) KINASES:5.6.3 PTK Group B membrane spanning protein tyrosine kinases.PTK XV Fibroblast growth factor receptor family .FGFR-3
	ec	ZA70_HUMAN	ZA70_HUMAN EC:2.7.1.112:TYROSINE-PROTEIN KINASE ZAP-70 (EC 2.7.1.112) (70 KDA ZETA-ASSOCIATED PROTEIN) (SYK-RELATED TYROSINE KINASE).
	Database	ID	Description
	scop	d1gjoa	d1gjoa_SCOP:d.144.1.2: Fibroblast growth factor receptor 2
	scop	d1ev2e1	d1ev2e1.SCOP:b.1.1.4: Fibroblast growth factor receptor, FGFR
	scop	d1gjoa	d1gjoa_SCOP:d.144.1.2: Fibroblast growth factor receptor 2

Protein Domains	scop	<u>d1ev2e1</u>	d1ev2e1 SCOP:b.1.1.4: Fibroblast growth factor receptor, FGFR	4.25E-21
	pfam	<u>ig</u>	Immunoglobulin domain	1.6E-5
	pfam	<u>ig</u>	Immunoglobulin domain	3.2E-8
	pfam	<u>pkinase</u>	Protein kinase domain	2.3E-92
	pfam	<u>ig</u>	Immunoglobulin domain	1.6E-5
	pfam	<u>ig</u>	Immunoglobulin domain	3.2E-8
	pfam	<u>pkinase</u>	Protein kinase domain	2.3E-92
	pfam	<u>ig</u>	Immunoglobulin domain	7.3E-8
	InterPro	IPR000719 <u>EMBL-EBI</u>	Protein kinase	
	InterPro	IPR007110 <u>EMBL-EBI</u>	Immunoglobulin-like	
Protein Domains	InterPro	IPR001245 <u>EMBL-EBI</u>	Tyrosine protein kinase	
	InterPro	IPR008266 <u>EMBL-EBI</u>	Tyrosine protein kinase, active site	
	InterPro	IPR003598 <u>EMBL-EBI</u>	Immunoglobulin C-2 type	

Trans Membrane

ID	Number Of Domains	Probability of Interior N-Terminus
NP_000133	2	0.11005

Sequence

>HUGENEFL:M64347_AT
gacttcaaagcaagctgggtattttcatataaaattcttctaatgctgtgtgtccaggca
gggagacggtttccaggaggggccggccctgtgtgcaggttccgatgttattagatgtt
acaagtttatatatatctatatataatttattgagttttacaagatgtatttgttgt
agacttaacacttcttacgcaatgcttctagagttttatagcctggactgtctacctttca
aagcttggagggaagccgtgaattcagttggttcgttctgtactgttactgggccctgag
tctgggcagctgtcccttgcttgccctgcagggccatggctcaggggtgggtctctcttggg
gccagtgcatgggtggccagaggtgtcacccaaaccggcaggtgcgatt

Target Sequence

Probe Info	Probe Sequence(5'-3')	Probe		Probe Interrogation Position	Strandedness
		X	Y		
Probe Info	GACTTCAAAGCAAGCTGGTATTTTC	359	161	3348	Antisense
	CATACAAATTCTTCTAATTGCTGTG	360	161	3372	Antisense
	AATTCTTCTAATTGCTGTGTGTCCC	361	161	3378	Antisense
	TGCTGTGTGTCCCAGGCAGGGAGAC	362	161	3390	Antisense
	TGTGTGCAGGTTCCGATGTTATTAG	363	161	3438	Antisense
	TCTTACGCAATGCTTCTAGAGTTTT	364	161	3540	Antisense
	GCAATGCTTCTAGAGTTTTATAGCC	365	161	3546	Antisense
	GAGTTTTATAGCCTGGACTGCTACC	366	161	3558	Antisense
	TGCTACCTTTCAAAGCTTGGAGGGA	367	161	3576	Antisense
	AAGCTTGGAGGGAAGCCGTGAATTC	368	161	3588	Antisense
	TGAATTCAGTTGGTTCTGTCTGTAC	369	161	3606	Antisense
	GTTCTGTCTGTACTGTTACTGGGCC	370	161	3618	Antisense
	CTGGGCCCTGAGTCTGGGCAGCTGT	371	161	3636	Antisense
	CCTGAGTCTGGGCAGCTGTCCCTTG	372	161	3642	Antisense
	TCTGGGCAGCTGTCCCTTGCTTGCC	373	161	3648	Antisense
	TCCCTTGCTTGCTGCAGGGCCATG	374	161	3660	Antisense

GCTTGCCTGCAGGGCCATGGCTCAG	375	161	3666	Antisense
CTTGGGGCCCAGTGCATGGTGGCCA	376	161	3702	Antisense
GTGGCCAGAGGTGTCACCCAAACCG	377	161	3720	Antisense
GTCACCCAAACCGGCAGGTGCGATT	378	161	3732	Antisense

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The Sequence Revision History tool allows you to see the various gi numbers, version numbers, and update dates for sequences that appeared in a specific GenBank record.

E.g., search for U46667 in the tool to see the old and current identifiers of the nucleotide sequence in that record.

Note that the original gi number for the nucleotide sequence, 2734632, does not have a corresponding version number. This is true because it was removed from the database (and replaced by 3172140) before the new accession.version system was implemented in Feb. 1999. At that time, each sequence in the GenBank/EMBL/DDBJ database received a version number of 1, even if they had been updated in the past.

In addition, if a GenBank record contains an updated sequence, the Comment field will contain a cross-reference to the gi number of the earlier sequence. (E.g., see U46667 in Entrez.) If you follow the link for that earlier gi number, Entrez will display that version of the GenBank record. Similarly, the Comment field of the older version will have a warning that the sequence has been updated, and will contain a cross-reference to the newer version.

More details about sequence identification numbers (GI and accession.version).

Back to sample record.

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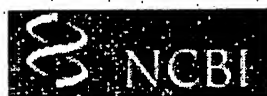
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Revised October 1, 2003

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Comments about site map to Renata Geer renata@ncbi.nlm.nih.gov



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GI	Version	Update Date	Status	I	II
3172140	1	Aug 7 1998 9:28 AM	Live	<input checked="" type="radio"/>	<input type="radio"/>
3172140	1	Jun 2 1998 4:31 PM	Dead	<input type="radio"/>	<input checked="" type="radio"/>
2734632	n/a	Jan 3 1998 12:12 AM	Dead	<input type="radio"/>	<input checked="" type="radio"/>
2734632	n/a	Jan 1 1998 12:30 AM	Dead	<input type="radio"/>	<input checked="" type="radio"/>

Accession U46667 was first seen at NCBI on Jan 1 1998 12:30 AM

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Details for HUGENEFL:L17131_RNA1_AT

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GeneChip Array Information

Probe Set ID L17131_ma1_at
GeneChip Array HumanGeneFL Array
Organism Common Name Human

Probe Design Information

Transcript ID L17131_ma1
Sequence Type Exemplar sequence
Representative Public ID L17131_NCB1
Target Description L17131, class A, 20 probes, 20 in L17131mRNA#1 1646-2198, Human high mobility group protein (HMG-I(Y)) gene exons 1-8, complete cds

Sequence

>HUGENEFL:L17131_RNA1_AT
ttgtccagggtgagggcccaagagccctgtggccgccacctgaggtgggctggggctgctcc
cctaaccctactttcgttccgccactcagccatttccccctcctcagatggggcaccaat
aacaaggagctcaccctgcccgtcccaacccccctcctgctcctccctgcccccaagg
ttctgggtccatttttctctgttcacaaactacctctggacagtgtgtgtgtttttgt
tcaatgttccattcttctgacatccgtcattgtgtgtgtaccagcgccaaatgttcattcc
tcattgacctctgttctgtcccacgatccctcccccaagatactcttgtggggaagagg
ggctggggcatggcaggctgggtgaccgactacccagtcaggaagggtggggccctg
ccctaggatgctgcagcagagtgagcaagggggcccgatcgaccataaagggtgtagg
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ctccctctctggtttctatttgcagttacttgaata

Target Sequence

Probe Sequence(5'-3')	Probe		Probe Interrogation Position	Strandedness
	X	Y		
TTGTCCAGGTGAGGCCCAAGAGCCC	294	101	1658	Antisense
AGGTGAGGCCCAAGAGCCCTGTGGC	295	101	1664	Antisense
ACCAATAACAAGGAGCTCACCCTGC	296	101	1772	Antisense
TTTTCCTCTGTTCAAACTACCTC	297	101	1850	Antisense
CTACCTCTGGACAGTTGTGTTGTTT	298	101	1868	Antisense
TTCCATTCTTCGACATCCGTCATTG	299	101	1904	Antisense
TCTTCGACATCCGTCATTGCTGCTG	300	101	1910	Antisense

Probe Info	GCTACCAGCGCCAAATGTTTCATCCT	301	101	1934	Antisense
	TCATCCTCATTGCCTCCTGTTCTGC	302	101	1952	Antisense
	TCATTGCCTCCTGTTCTGCCCACGA	303	101	1958	Antisense
	AAGATACTCTTTGTGGGGAAGAGGG	304	101	1994	Antisense
	GCAGGCTGGGTGACCGACTACCCCA	305	101	2030	Antisense
	CCCCTAGGATGCTGCAGCAGAGTGA	306	101	2078	Antisense
	AGCAAGGGGGCCCGAATCGACCATA	307	101	2102	Antisense
	CGAATCGACCATAAAGGGTGTAGGG	308	101	2114	Antisense
	GCCATGATTTGTCCCAGCCTGGGGC	309	101	2174	Antisense
	CTGGGGCTCCCTCTCTGGTTTCCTA	310	101	2192	Antisense
	CTCCCTCTCTGGTTTCCTATTTGCA	311	101	2198	Antisense
	CTCTGGTTTCCTATTTGCAGTTACT	312	101	2204	Antisense
	TTTCCTATTTGCAGTTACTTGAATA	313	101	2210	Antisense

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- (0) All Descriptions (L17131_rnal_at)
- (1) All Descriptions (m64347)

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GeneChip Array Information

Probe Set ID X74801_at
 GeneChip Array HumanGeneFL Array
 Organism Common Name Human

Probe Design Information

Transcript ID X74801
 Sequence Type Exemplar sequence
 Representative Public ID X74801 [NCBI](#)
 Target Description X74801, class B, 20 probes, 12 in X74801cds 1282-1552: 8 in reverseSequence, 1636-1837, H.sapiens Cctg mRNA for chaperonin

Genomic Alignment of Target Sequence

Assembly April 2003 (NCBI 33)
 Alignment(s)

	Position	% Identity	Cytoband
chr1: 153495555-153497649 (-)	UCSC	100	q22

Overlapping Transcripts	Representative Transcript	UniGene Description	Position
	NM_005998 NCBI	chaperonin containing TCP1, subunit 3 (gamma)	chr1:153495551-153524840 (-) UCSC

Public Domain and Genome References

Gene Title chaperonin containing TCP1, subunit 3 (gamma)
 Gene Symbol CCT3 [HGNC](#)
 Chromosomal Location 1q23
 UniGene ID Hs.1708 [NCBI](#) (FULL LENGTH)
 Ensembl ENSG00000163468 [Ensembl](#)
 LocusLink 7203 [NCBI](#)
 SwissProt AAH06501 [EMBL-EBI](#)
 P49368 [EMBL-EBI](#)
 OMIM 600114 [NCBI](#)
 RefSeq Protein ID NP_005989 [NCBI](#)

RefSeq	RefSeq Transcript ID	RefSeq Title
	NM_005998 <u>NCBI</u>	chaperonin containing TCP1, subunit 3 (gamma)
Functional Annotations		
	ID	Title Organism Type
	<u>ATH1-121501:246830 AT</u>	chaperonin, putative Arabidopsis Putative Ortholog
	<u>ATGENOME1:18906 AT</u>	chaperonin, putative Arabidopsis Putative Ortholog
	<u>DROSGENOME1:153982 AT</u>	Drosophila Putative Ortholog
	<u>MG-U74AV2:161238 F AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MG-U74AV2:98153 AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MG-U74CV2:171548 AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MOE430A:1416024 X AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MOE430A:1426067 X AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MOE430A:1448178 A AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MOE430A:1449645 S AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MOE430A:1451915 AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MOE430A:1459987 S AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
Ortholog	<u>MU11KSUBA:C79428 RC F AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MU11KSUBA:L20509 F AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MOUSE430 2:1416024 X AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MOUSE430 2:1426067 X AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MOUSE430 2:1448178 A AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MOUSE430 2:1449645 S AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MOUSE430 2:1451915 AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MOUSE430 2:1459987 S AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MOUSE430A 2:1416024 X AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MOUSE430A 2:1426067 X AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MOUSE430A 2:1448178 A AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MOUSE430A 2:1449645 S AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog
	<u>MOUSE430A 2:1451915 AT</u>	chaperonin subunit 3 (gamma) Mouse Curated Ortholog

MOUSE430A 2:1459987 S AT chaperonin subunit 3 Mouse (gamma) **Curated Ortholog**

GO Biological Process (view graph)

ID	Description	Evidence	Links
6457	protein folding	traceable author statement	QuickGO AmiGO

GO Cellular Component (view graph)

ID	Description	Evidence	Links
5829	cytosol	not recorded	QuickGO AmiGO
5856	cytoskeleton	traceable author statement	QuickGO AmiGO

GO Molecular Function (view graph)

ID	Description	Evidence	Links
3754	chaperone activity	traceable author statement	QuickGO AmiGO
5524	ATP binding	inferred from electronic annotation	QuickGO AmiGO

Method	ID	Description	E-Value
blast	33873532		0.0
blast	31542292	chaperonin containing TCP1, subunit 3 (gamma); TCP1 (t-complex-1) ring complex, polypeptide 5 [Homo sapiens]	0.0

Database	ID	Description	E-Value
scop	d1a6da3	d1a6da3 SCOP:d.56.1.2; Thermosome	4.08E-25
scop	d1gmla	d1gmla_ SCOP:c.8.5.2; Thermosome	1.01E-57
scop	d1a6da1	d1a6da1 SCOP:a.129.1.2; Thermosome	4.81E-83
pfam	cpn60_TCP1	TCP-1/cpn60 chaperonin family	5.7E-210
InterPro	IPR002423	Chaperonin Cpn60/TCP-1	
InterPro	IPR001844	Chaperonin Cpn60	
InterPro	IPR002194	Chaperonin TCP-1	
InterPro	IPR008950	GroEL-like chaperone, ATPase	

Sequence

Target Sequence

>HUGENEFL: X74801_AT
 atgactggtgtggaacaatggccatacagggtctgttgccaggccctagaggtcattcct
 cgtaccctgatccagaactgtggggccagcaccatccgtctacttacctcccttggggcc
 aagcacaccaggagaactgtgagacctggggtgtaaatggtgagacgggtactttgggtg
 gacatgaaggaaactgggcataatgggagccattggctgtgaagctgcagactataagaca
 gcagtggagacggcagttctgtactgcgaattgatgacatcgtttcaggccacaaaaag
 aaaggcgatgaccagagccggcaaggcgggctcctgatgctggccaggagtgaagtgc
 ggcaaggctacttcaatgcacagaaccagcagagtctcccttttcttgagccagagtgc
 caggaacactgtggacgtctttgttcagaagggatcaggttggggggcagccccagtc
 ctttctgtcccagctcagttttccaaaagacactgacatgtaattcttctctattgtaag
 gtttccatttagtttgctccgatgattaaatctaagtca

Probe Sequence(5'-3')	Probe X	Probe Y	Probe Interrogation Position	Strandedness
-----------------------	---------	---------	------------------------------	--------------

Probe Info	ATGACTGGTGTGGAACAATGGCCAT	60	345	1294	Antisense
	GAACAATGGCCATACAGGGCTGTTG	61	345	1306	Antisense
	CTGATCCAGAACTGTGGGGCCAGCA	62	345	1360	Antisense
	CAGAACTGTGGGGCCAGCACCATCC	63	345	1366	Antisense
	TGTGGGGCCAGCACCATCCGTCTAC	64	345	1372	Antisense
	CTGGGCATATGGGAGCCATTGGCTG	65	345	1486	Antisense
	ATATGGGAGCCATTGGCTGTGAAGC	66	345	1492	Antisense
	GAGCCATTGGCTGTGAAGCTGCAGA	67	345	1498	Antisense
	TTGGCTGTGAAGCTGCAGACTTATA	68	345	1504	Antisense
	GAGACGGCAGTTCTGCTACTGCGAA	69	345	1540	Antisense
	GCAGTTCTGCTACTGCGAATTGATG	70	345	1546	Antisense
	ATTGATGACATCGTTTCAGGCCACA	71	345	1564	Antisense
	GTGCTAGGCAAGGCTACTTCAATGC	72	345	1648	Antisense
	GGCAAGGCTACTTCAATGCACAGAA	73	345	1654	Antisense
	GCTACTTCAATGCACAGAACCAGCA	74	345	1660	Antisense
	CACAGAACCAGCAGAGTCTCCCCTT	75	345	1672	Antisense
	GAGCCAGAGTGCCAGGAACACTGTG	76	345	1702	Antisense
	CACTGACATGTAATTCTTCTCTATT	77	345	1804	Antisense
	TAGTTTGCTTCCGATGATTAAATCT	78	345	1843	Antisense
	GCTTCCGATGATTAAATCTAAGTCA	79	345	1849	Antisense



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Details for HUGENEFL:U15008_AT

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NetAffx Links: [Cluster Members](#)
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GeneChip Array Information

Probe Set ID U15008_at
GeneChip Array HumanGeneFL Array
Organism Common Name Human

Probe Design Information

Transcript ID U15008
Sequence Type Exemplar sequence
Representative Public ID U15008 [NCBI](#)
Target Description U15008, class A, 20 probes, 20 in U15008 25-433, Human SnRNP core protein Sm D2 mRNA, complete cds

Genomic Alignment of Target Sequence

Assembly April 2003 (NCBI 33)
Alignment(s) Position % Identity Cytoband
chr19: 50882580-50883664 (-) [UCSC](#) 98 q13.32

Overlapping Transcripts	Representative Transcript	UniGene Description	Position
	NM_004597 NCBI	small nuclear ribonucleoprotein D2 polypeptide 16.5kDa	chr19:50882558-50887282 (-) UCSC
	NM_177542 NCBI	small nuclear ribonucleoprotein D2 polypeptide 16.5kDa	chr19:50882558-50887282 (-) UCSC

Public Domain and Genome References

Gene Title small nuclear ribonucleoprotein D2 polypeptide 16.5kDa
Gene Symbol SNRPD2 [HGNC](#)
Chromosomal Location 19q13.2
UniGene ID Hs.424327 [NCBI](#) (FULL LENGTH)
Ensembl ENSG00000125743 [Ensembl](#)
LocusLink 6633 [NCBI](#)
SwissProt P43330 [EMBL-EBI](#)
OMIM 601061 [NCBI](#)

RefSeq Protein ID	NP_004588 NCBI
	NP_808210 NCBI
RefSeq Transcript ID	
RefSeq	NM_004597 NCBI small nuclear ribonucleoprotein polypeptide D2
	NM_177542 NCBI small nuclear ribonucleoprotein polypeptide D2

Functional Annotations

	ID	Title	Organism	Type
Ortholog	ATH1-121501:266482_AT	small nuclear ribonucleo protein D2 -related	Arabidopsis	Putative Ortholog
	C. ELEGANS:172931_X_AT	small nuclear ribonucleoprotein D2 like	Celegans	Putative Ortholog
	DROSGENOME1:153483_AT		Drosophila	Putative Ortholog
	MG-U74AV2:95049_AT	small nuclear ribonucleoprotein D2	Mouse	Curated Ortholog
	MOE430A:1452680_AT	small nuclear ribonucleoprotein D2	Mouse	Curated Ortholog
	MU11KSUBA:AA271024_S_AT	small nuclear ribonucleoprotein D2	Mouse	Curated Ortholog
	MOUSE430_2:1452680_AT	small nuclear ribonucleoprotein D2	Mouse	Curated Ortholog
	MOUSE430A_2:1452680_AT	small nuclear ribonucleoprotein D2	Mouse	Curated Ortholog

GO Biological Process (view graph)

ID	Description	Evidence	Links
245	spliceosome assembly	traceable author statement	QuickGO AmiGO
6371	mRNA splicing	traceable author statement	QuickGO AmiGO

GO Cellular Component (view graph)

	ID	Description	Evidence	Links
Gene Ontology	5681	spliceosome complex	traceable author statement	QuickGO AmiGO
	5732	small nucleolar ribonucleoprotein complex	inferred from electronic annotation	QuickGO AmiGO
	30532	small nuclear ribonucleoprotein complex	traceable author statement	QuickGO AmiGO

GO Molecular Function (view graph)

ID	Description	Evidence	Links
8248	pre-mRNA splicing factor activity	inferred from electronic annotation	QuickGO AmiGO

Protein Similarities

Method	ID	Description	E-Value
blast	4759158	small nuclear ribonucleoprotein polypeptide D2; snRNP core protein D2 [Homo sapiens]	1.0E-62
blast	26337731		3.0E-62
blast	4759158	small nuclear ribonucleoprotein polypeptide D2; snRNP core protein D2 [Homo sapiens]	1.0E-62
blast	26337731		3.0E-62

Database	ID	Description	E-Value
scop	d1b34b	d1b34b SCOP:b.38.1.1.; D2 core SNRNP	1.85E-

Protein Domains	scop	d1b34b	protein d1b34b_SCOP:b.38.1.1: D2 core SNRNP protein	28 1.85E-28
	pfam	LSM	LSM domain	1.1E-16
	pfam	LSM	LSM domain	1.1E-16
	InterPro	IPR001163	Small nuclear ribonucleoprotein (Sm protein)	
		EMBL-EBI		

Sequence

Target Sequence

>HUGENEFL:U15008_AT
 accatcatgagcctcctcaacaagcccaagagtgagatgacccagaggagctgcagaag
 cgagaggaggaggaatttaacaccgggccactctctgtgctcacacagtcagtcagaac
 aatacccaagtgtcatcaactgccgcaacaataagaaactcctgggccgctgaaggcc
 ttcgataggcactgcaacatggtgctggagaacgtgaaggagatgtggactgaggtaccc
 aagagtggcaagggcaagaagaagtcgaagccagtcacaaagaccgctacatctccaag
 atgttctgctgcggggactcagtcacgtggtcctgcccgaaccgctcatgcggccaag
 taggggcccgtgtctgttgacagaactcactcctctgtcctatgaagaccgtgccatt
 ggtgttgagaata

Probe Info	Probe Sequence(5'-3')	Probe		Probe Interrogation Position	Strandedness
		X	Y		
	ACCATCATGAGCCTCCTCAACAAGC	99	211	37	Antisense
	AGTGAGATGACCCAGAGGAGCTGC	100	211	67	Antisense
	AACACCGGTCCACTCTCTGTGCTCA	101	211	115	Antisense
	GGTCCACTCTCTGTGCTCACACAGT	102	211	121	Antisense
	CTCTCTGTGCTCACACAGTCAGTCA	103	211	127	Antisense
	GTGCTCACACAGTCAGTCAAGAACA	104	211	133	Antisense
	TCAGTCAAGAACAATACCCAAGTGC	105	211	145	Antisense
	AATACCCAAGTGCTCATCAACTGCC	106	211	157	Antisense
	CAAGTGCTCATCAACTGCCGCAACA	107	211	163	Antisense
	CGCGTGAAGGCCCTTCGATAGGCACT	108	211	205	Antisense
	AAGGCCTTCGATAGGCACTGCAACA	109	211	211	Antisense
	TTCGATAGGCACTGCAACATGGTGC	110	211	217	Antisense
	GTACCCAAGAGTGGCAAGGGCAAGA	111	211	271	Antisense
	TACATCTCCAAGATGTTCTGCGCG	112	211	325	Antisense
	TCAGTCATCGTGGTCCTGCGGAACC	113	211	355	Antisense
	TAGGGGCCGCCTGTCTGTTGACAGA	114	211	397	Antisense
	TGACAGAACTCACTCCTCTGTCCTA	115	211	415	Antisense
	CTCCTCTGTCCTATGAAGACCGCTG	116	211	427	Antisense
	TGTCCTATGAAGACCGCTGCCATTG	117	211	433	Antisense
	ACCGCTGCCATTGGTGTGAGAATA	118	211	445	Antisense

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Details for HUGENEFL:AFFX-BIOB-M_ST

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GeneChip Array Information

Probe Set ID AFFX-BioB-M_st
**GeneChip
Array** HumanGeneFL Array
**Organism
Common
Name** Human

Probe Design Information

Transcript ID AFFX-BioB-M
**Sequence
Type** Control sequence
**Representative
Public ID** J04423 [NCBI](#)
**Target
Description** J04423 E coli bioB gene biotin synthetase (-5, -M, -3 represent transcript regions
5 prime, Middle, and 3 prime respectively)

Sequence

Target Sequence

```
>HUGENEFL:AFFX-BIOB-M_ST
gccggagttttacggcaatatcatcaccacgcacattatcaggaacgcctcgatacgct
ggaaaaagtgcgcgatgccgggatcaaagtctgttctggcggcattgtgggcttaggcga
aacggtaaaagatcgcccggtattattgtgcaactggcaaacctgccgacgcccgga
aagcgtgccaatcaacatgctggtgaagtgaaaggcacgccgcttgccgataacgatga
tgtcgatgcctttgattt
```

Probe Info

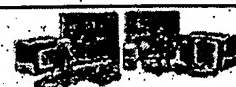
Probe Sequence(5'-3')	Probe		Probe Interrogation Position	Strandedness
	X	Y		
GATGATATTGCCGTA AAACTCCGGC	201	11	483	Sense
TGTGGTGATGATATTGCCGTA AAAC	202	11	489	Sense
TAAGTGCGTGTTGGTGATGATATTGC	203	11	497	Sense
GTTCTTGATAAGTGCGTGTTGGTGAT	204	11	505	Sense
ATCGAGGCGTTCCCTGATAAGTGCGT	205	11	513	Sense
GCATCGCGCACTTTTCCAGCGTAT	206	11	536	Sense
GATCCCGGCATCGCGCACTTTTCC	207	11	543	Sense
GACTTTGATCCCGGCATCGCGCACT	208	11	549	Sense
CGCCAGAACAGACTTTGATCCCGGC	209	11	559	Sense
CCCACAATGCCGCCAGAACAGACTT	210	11	569	Sense

TGCAGCAATAATCCGGCGCGATCTT	211	11	611	Sense
TTGCCAGTTGCAGCAATAATCCGGC	212	11	619	Sense
CGGCAGGTTTGCCAGTTGCAGCAAT	213	11	627	Sense
ATGTTGATTGGCACGCTTTCGGGCG	214	11	656	Sense
CACCAGCATGTTGATTGGCACGCTT	215	11	663	Sense
TTCACCTTCACCAGCATGTTGATTG	216	11	671	Sense
AGCGGCGTGCCTTTCACCTTCACCA	217	11	683	Sense
CATCATCGTTATCGGCAAGCGGCGT	218	11	700	Sense
GCATCGACATCATCGTTATCGGCAA	219	11	707	Sense
AAATCAAAGGCATCGACATCATCGT	220	11	716	Sense

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→ (2) All Descriptions
(AFFX-BioDn-5)

→ (2) All Descriptions
(AFFX-BioB-M)

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Details for HUGENEFL:AFFX-BIODN-5_ST

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GeneChip Array Information

Probe Set ID AFFX-BioDn-5_st
GeneChip Array HumanGeneFL Array
Organism Common Name Human

Probe Design Information

Transcript ID AFFX-BioDn-5
Sequence Type Control sequence
Representative Public ID J04423 [NCBI](#)
Target Description J04423 E coli bioD gene dethiobiotin synthetase (-5 and -3 represent transcript regions 5 prime and 3 prime respectively)

Sequence

Target
Sequence

>HUGENEFL:AFFX-BIODN-5_ST
gggaaaaactgtcgccagttgtgcacttttacaagccgcaaaaggcagcaggctaccggacg
gcaggttataaaacgggtcgccctctggcagcgaaaagaccccggaaggtttacgcaatagc
gacgcgctggcgttacagcgcaacagcagcctgcagctggattacgcaacagtaaatcct
tacaccttcgcagaaccacttcgcgcacatcatcagcgcgcaagagggcagaccgata
gaatcattggtaatgagcgccggattacgcgcgcttg

Probe Info

Probe Sequence(5'-3')	Probe		Probe Interrogation Position	Strandedness
	X	Y		
GTGCACAACTGGCGACAGTTTTCCC	281	11	49	Sense
GGCTTGTA AAAAGTGCACAACTGGCG	282	11	60	Sense
GCTGCCCTTTGCGGCTTGTA AAAAGTG	283	11	71	Sense
GGTAGCCTGCTGCCTTTGCGGCTTG	284	11	79	Sense
CCGTCCGGTAGCCTGCTGCCTTTGC	285	11	85	Sense
CAGCGCGTCGCTATTGCGTAAACCT	286	11	153	Sense
GTAACGCCAGCGCGTCGCTATTGCG	287	11	160	Sense
TTGCGCTGTAACGCCAGCGCGTCGC	288	11	167	Sense
TGCTGTTGCGCTGTAACGCCAGCGC	289	11	172	Sense
TGCAGGCTGCTGTTGCGCTGTAACG	290	11	179	Sense

TCCAGCTGCAGGCTGCTGTTGCGCT	291	11	185	Sense
TGCGTAATCCAGCTGCAGGCTGCTG	292	11	192	Sense
TTACTGTTGCGTAATCCAGCTGCAG	293	11	199	Sense
CGGTCTGCCCTCTTGCGCGCTGATG	294	11	261	Sense
GATTCTATCGGTCTGCCCTCTTGCG	295	11	269	Sense
TACCAATGATTCTATCGGTCTGCCC	296	11	276	Sense
CTCATTACCAATGATTCTATCGGTC	297	11	281	Sense
TCCGGCGCTCATTACCAATGATTCT	298	11	288	Sense
CGCGTAATCCGGCGCTCATTACCAA	299	11	295	Sense
CAAGCGCGCGTAATCCGGCGCTCAT	300	11	301	Sense

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Details for HUGENEFL:X15880_AT

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GeneChip Array Information

Probe Set ID X15880_at
GeneChip Array HumanGeneFL Array
Organism Common Name Human

Probe Design Information

Transcript ID X15880
Sequence Type Exemplar sequence
Representative Public ID X15880 [NCBI](#)
Target Description X15880, class C, 20 probes, 20 in all_X15880 1690-2273, Human mRNA for collagen VI alpha-1 C-terminal globular domain

Genomic Alignment of Target Sequence

Assembly	Position	% Identity	Cytoband
April 2003 (NCBI 33)	chr21: 46280561-46281145 (+) UCSC	100	q22.3

Overlapping Transcripts	Representative Transcript	UniGene Description	Position
	NM_001848 NCBI	collagen, type VI, alpha 1	chr21:46257869-46281164 (+) UCSC

Public Domain and Genome References

Gene Title collagen, type VI, alpha 1
Gene Symbol COL6A1 [HGNC](#)
Chromosomal Location 21q22.3
UniGene ID Hs.415997 [NCBI](#) (FULL LENGTH)
Ensembl ENSG00000142156 [Ensembl](#)
LocusLink 1291 [NCBI](#)
SwissProt P12109 [EMBL-EBI](#)
Q7Z645 [EMBL-EBI](#)
Q8TBN2 [EMBL-EBI](#)
Q9BSA8 [EMBL-EBI](#)
OMIM 120220 [NCBI](#)

RefSeq Protein ID NP_001839 [NCBI](#)

RefSeq RefSeq Transcript ID RefSeq Title
 NM_001848 [NCBI](#) collagen, type VI, alpha 1 precursor

Functional Annotations

	ID	Title	Organism	Type
Ortholog	MG-U74AV2:162459 F AT	procollagen, type VI, alpha 1	Mouse	Curated Ortholog
	MG-U74AV2:95493 AT	procollagen, type VI, alpha 1	Mouse	Curated Ortholog
	MOE430A:1448590 AT	procollagen, type VI, alpha 1	Mouse	Curated Ortholog
	MU11KSUBB:X66405 S AT	procollagen, type VI, alpha 1	Mouse	Curated Ortholog
	MOUSE430 2:1448590 AT	procollagen, type VI, alpha 1	Mouse	Curated Ortholog
	MOUSE430A 2:1448590 AT	procollagen, type VI, alpha 1	Mouse	Curated Ortholog

GO Biological Process (view graph)

ID	Description	Evidence	Links
7155	cell adhesion	non-traceable author statement	QuickGO AmiGO

GO Cellular Component (view graph)

ID	Description	Evidence	Links
5578	extracellular matrix	inferred from electronic annotation	QuickGO AmiGO
5589	collagen type VI	non-traceable author statement	QuickGO AmiGO

Gene Ontology

GO Molecular Function (view graph)

ID	Description	Evidence	Links
5194	cell adhesion molecule activity	inferred from electronic annotation	QuickGO AmiGO
5201	extracellular matrix structural constituent	inferred from electronic annotation	QuickGO AmiGO

Protein Similarities	Method	ID	Description	E-Value
	blast	15011913		0.0
	blast	13878903		0.0

Protein Domains	Database	ID	Description	E-Value
	scop	d1atza	d1atza_SCOP:c.62.1.1:] von Willebrand factor A3 domain	3.63E-37
	pfam	vwa	von Willebrand factor type A domain	9.6E-24
	pfam	vwa	von Willebrand factor type A domain	4.7E-32
	pfam	vwa	von Willebrand factor type A domain	2.7E-35
	pfam	Collagen	Collagen triple helix repeat (20 copies)	2.4E-11
	pfam	Collagen	Collagen triple helix repeat (20 copies)	3.8E-14
	pfam	Collagen	Collagen triple helix repeat (20 copies)	3.3E-10
	pfam	Collagen	Collagen triple helix repeat (20 copies)	2.6E-11
	InterPro	IPR008161 EMBL-EBI	Collagen helix repeat	
	InterPro	IPR002035 EMBL-EBI	von Willebrand factor, type A	

InterPro IPR008160 Collagen triple helix repeat
EMBL-EBI

Sequence

>HUGENEFL:X15880_AT

Target
Sequence

agcaagacgcctctcggggcctgtgcccactagcctccctctcctctgtcccatagct
ggtttttccaccaatcctcacctaacagttactttacaattaaactcaaagcaagctct
tctcctcagcttggggcagccattggcctctgtctcgtttgggaaaccaaggtcaggag
gccgttgacacataaatctcggcgactcgccccgtctcctgaggggtcctgctggtagc
cggcctggaccttggccctacagccctggaggccgctgctgaccagcactgaccccgacc
tcagagagtactcgcagggcgctggctgcactcaagaccctcgagattaacggtgctaa
ccccgtctgctctccctcccgcagagactggggcctggactggacatgagagccccttg
gtgccacagagggctgtgtcttactagaaacaacgcaaacctctccttctcagaatagt
gatgtgttcgacgttttatcaaaggccccctttctatgttcattgttagttttgctccttc
tgtgttttttctgaaccatatccatgttgctgacttttccaa

Probe Sequence(5'-3')	Probe		Probe	Strandedness
	X	Y	Interrogation Position	
AGCAAGACGCCTCTCGGGGCCTGTG	76	317	1702	Antisense
AAACTCAAAGCAAGCTCTTCTCCTC	77	317	1804	Antisense
AAAGCAAGCTCTTCTCCTCAGCTTG	78	317	1810	Antisense
TCTCCTCAGCTTGGGGCAGCCATTG	79	317	1822	Antisense
GCCATTGGCCTCTGTCTCGTTTTGG	80	317	1840	Antisense
GCAGACATAAATCTCGGCGACTCGG	81	317	1888	Antisense
GCCCCGTCTCCTGAGGGTCTGCTG	82	317	1912	Antisense
TGGCCCTACAGCCCTGGAGGCCGCT	83	317	1954	Antisense
TCAGAGAGTACTCGCAGGGGCGCTG	84	317	2002	Antisense
AGTACTCGCAGGGGCGCTGGCTGCA	85	317	2008	Antisense
GGCGCTGGCTGCACTCAAGACCCTC	86	317	2020	Antisense
GGACATGAGAGCCCCCTTGGTGCCAC	87	317	2104	Antisense
GAGAGCCCCCTTGGTGCCACAGAGGG	88	317	2110	Antisense
CCCTTGGTGCCACAGAGGGCTGTGT	89	317	2116	Antisense
GTGCCACAGAGGGCTGTGTCTTACT	90	317	2122	Antisense
CAGAGGGCTGTGTCTTACTAGAAAC	91	317	2128	Antisense
CTCCTTCCTCAGAATAGTGATGTGT	92	317	2164	Antisense
TTTTTCTGAACCATATCCATGTTGC	93	317	2248	Antisense
TGAACCATATCCATGTTGCTGACTT	94	317	2254	Antisense
ATATCCATGTTGCTGACTTTTCCAA	95	317	2260	Antisense

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- [-> all probe sets \(7129\)](#)

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Details for HUGENEFL:U23752_AT

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GeneChip Array Information

Probe Set ID U23752_at
 GeneChip Array HumanGeneFL Array
 Organism Common Name Human

Probe Design Information

Transcript ID U23752
 Sequence Type Exemplar sequence
 Representative Public ID U23752 [NCBI](#)
 Target Description U23752, class A, 20 probes, 20 in U23752 1679-1919, Human SOX-11 mRNA, complete cds

Genomic Alignment of Target Sequence

Assembly April 2003 (NCBI 33)
 Alignment(s) Position % Identity Cytoband
 chr2: 5856192-5856457 (+) [UCSC](#) 99 p25.2

Overlapping Transcripts	Representative Transcript	UniGene Description	Position
	NM_003108 NCBI	SRY (sex determining region Y)-box 11	chr2:5854537-5863255 (+) UCSC

Public Domain and Genome References

Gene Title SRY (sex determining region Y)-box 11
 Gene Symbol SOX11 [HGNC](#)
 Chromosomal Location 2p25
 UniGene ID Hs.432638 [NCBI](#) (FULL LENGTH)
 Ensembl ENSG00000176887 [Ensembl](#)
 LocusLink 6664 [NCBI](#)
 SwissProt P35716 [EMBL-EBI](#)
 OMIM 600898 [NCBI](#)
 RefSeq Protein ID NP_003099 [NCBI](#)

RefSeq Transcript ID RefSeq Title
 NM_003108 [NCBI](#) SRY-box 11

Functional Annotations

	ID	Title	Organism	Type
Ortholog	RAE230A:1387275_AT	SRY-box containing gene	Rat	Putative Ortholog
	RG-U34A:AJ004858_AT	SRY-box containing gene	Rat	Putative Ortholog

GO Biological Process (view graph)

ID	Description	Evidence	Links
6355	regulation of transcription, DNA-dependent	inferred from electronic annotation	QuickGO AmiGO
7399	neurogenesis	traceable author statement	QuickGO AmiGO

Gene Ontology

GO Cellular Component (view graph)

ID	Description	Evidence	Links
5634	nucleus	inferred from electronic annotation	QuickGO AmiGO

GO Molecular Function (view graph)

ID	Description	Evidence	Links
3677	DNA binding	inferred from electronic annotation	QuickGO AmiGO

	Method	ID	Description	E-Value
Protein Similarities	blast	4507161	SRY-box 11; SRY (sex-determining region Y)-box 11; SRY-related HMG-box gene 11; transcription factor SOX-11 [Homo sapiens]	0.0
	blast	23831472		0.0

	Database	ID	Description	E-Value
Protein Domains	scop	d1i11a	d1i11a_SCOP:a.21.1.1: Sox-5	2.36E-19
	pfam	HMG_box	HMG (high mobility group) box	1.1E-33
	InterPro	IPR000910	HMG1/2 (high mobility group) box	
		EMBL-EBI		

Sequence

Target Sequence

>HUGENEFL:U23752_AT
 ctccctttatcgtgtctcaaggtagttgcatacctagtctggagttgtgattatttccc
 aaaaaatgtgtttttgtaattactatttcttttccctgaaattcgtgattgcaacaaagg
 cagagggggcggcgcggcgaggaggtaggaccgcctccgaaggcgctgttgaagc
 ttgtcggctcttgaagtctggaagacgtctgcagaggacctttggcagcacaactgtt
 actctaggaggttggtggagatatt

Probe Sequence(5'-3')	Probe-Probe		Probe Interrogation Position	Strandedness
	X	Y		
CTTCCTTTATCGTGTCTCAAGGTAG	503	219	1691	Antisense
TTATCGTGTCTCAAGGTAGTTGCAT	504	219	1697	Antisense
TCGTGTCTCAAGGTAGTTGCATACC	505	219	1700	Antisense
AAGGTAGTTGCATACCTAGTCTGGA	506	219	1709	Antisense
GTAGTTGCATACCTAGTCTGGAGTT	507	219	1712	Antisense

Probe Info	GTTGCATACCTAGTCTGGAGTTGTG	508	219	1715	Antisense
	TACCTAGTCTGGAGTTGTGATTATT	509	219	1721	Antisense
	CTAGTCTGGAGTTGTGATTATTTTC	510	219	1724	Antisense
	TGTGATTATTTTCCCAAAAATGTG	511	219	1736	Antisense
	TTTTCCTGAAATTCGTGATTGCAAC	512	219	1781	Antisense
	GCTCCGGAAGGCGCTGTTTGAAGCT	513	219	1847	Antisense
	GCTGTTTGAAGCTTGTCGGTCTTTG	514	219	1859	Antisense
	TGAAGCTTGTCGGTCTTTGAAGTCT	515	219	1865	Antisense
	TTGTCGGTCTTTGAAGTCTGGAAGA	516	219	1871	Antisense
	TGGAAGACGTCTGCAGAGGACCCTT	517	219	1889	Antisense
	AAGACGTCTGCAGAGGACCCTTTTG	518	219	1892	Antisense
	GCAGAGGACCCTTTTGGCAGCACAA	519	219	1901	Antisense
	AGCACAACTGTTACTCTAGGGAGTT	520	219	1919	Antisense
	ACTGTTACTCTAGGGAGTTGGTGGA	521	219	1925	Antisense
	ACTCTAGGGAGTTGGTGGAGATATT	522	219	1931	Antisense

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Probe Set ID M12625_at
GeneChip Array HumanGeneFL Array
Organism Common Name Human

Probe Design Information
Transcript ID M12625
Sequence Type Exemplar sequence
Representative Public ID M12625 [NCBI](#)
Target Description M12625, class B, 20 probes; 13 in M12625mRNA 893-1259: 7 in reverseSequence, 1599-1683, Human lecithin-cholesterol acyltransferase mRNA, complete cds, with 5' and 3' flanking DNA sequences

Genomic Alignment of Target Sequence
Assembly April 2003 (NCBI 33)
Alignment(s)

	Position	% Identity	Cytoband
chr16: 67749925-67750484 (-)	UCSC	100	q22.1

Overlapping Transcripts	Representative Transcript	UniGene Description	Position
M12625 NCBI	lecithin-cholesterol acyltransferase	chr16:67749888-67754507 (-)	UCSC

Public Domain and Genome References
Gene Title lecithin-cholesterol acyltransferase
Gene Symbol LCAT [HGNC](#)
Chromosomal Location 16q22.1
UniGene ID Hs.387239 [NCBI](#) (FULL LENGTH)
Ensembl ENSG00000103080 [Ensembl](#)
LocusLink 3931 [NCBI](#)
SwissProt AAP88750 [EMBL-EBI](#)
P04180 [EMBL-EBI](#)
EC 2.3.1.43
OMIM 606967 [NCBI](#)

RefSeq Protein ID NP_000220 NCBI

RefSeq RefSeq Transcript ID RefSeq Title
NM_000229 NCBI lecithin-cholesterol acyltransferase precursor

Functional Annotations

	ID	Title	Organism	Type
Ortholog	MG-U74AV2:103023 AT	lecithin cholesterol acyltransferase	Mouse	Curated Ortholog
	MG-U74AV2:161759 R AT	lecithin cholesterol acyltransferase	Mouse	Curated Ortholog
	MOE430A:1417043 AT	lecithin cholesterol acyltransferase	Mouse	Curated Ortholog
	MU11KSUBA:J05154 S AT	lecithin cholesterol acyltransferase	Mouse	Curated Ortholog
	RAE230A:1367887 AT	lecithin cholesterol acyltransferase	Rat	Curated Ortholog
	RG-U34A:X54096 AT	lecithin cholesterol acyltransferase	Rat	Curated Ortholog
	MOUSE430 2:1417043 AT	lecithin cholesterol acyltransferase	Mouse	Curated Ortholog
	MOUSE430A 2:1417043 AT	lecithin cholesterol acyltransferase	Mouse	Curated Ortholog

GO Biological Process (view graph)

ID	Description	Evidence	Links
6629	lipid metabolism	inferred from electronic annotation	QuickGO AmiGO

GO Cellular Component (view graph)

ID	Description	Evidence	Links
5576	extracellular	not recorded	QuickGO AmiGO

Gene Ontology

GO Molecular Function (view graph)

ID	Description	Evidence	Links
4607	phosphatidylcholine-sterol O-acyltransferase activity	inferred from electronic annotation	QuickGO AmiGO
8415	acyltransferase activity	inferred from electronic annotation	QuickGO AmiGO
16740	transferase activity	inferred from electronic annotation	QuickGO AmiGO

Protein Similarities

Method	ID	Description	E-Value
blast	32879837		0.0
blast	4557892	lecithin-cholesterol acyltransferase precursor [Homo sapiens]	0.0

Protein Families

Method	ID	Description	E-Value
ec	LCAT_HUMAN	LCAT_HUMAN EC:2.3.1.43:PHOSPHATIDYLCHOLINE-STEROL ACYLTRANSFERASE PRECURSOR (EC 2.3.1.43) (LECITHIN-CHOLESTEROL ACYLTRANSFERASE) (PHOSPHOLIPID-CHOLESTEROL ACYLTRANSFERASE).	1.85E-171

Database	ID	Description	E-Value
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scop d1tca d1tca__ SCOP:c.69.1.17:| Triacylglycerol lipase 5.3E-8
 pfam LACT Lecithin:cholesterol acyltransferase 1.7E-182

InterPro IPR003386 Lecithin:cholesterol acyltransferase
EMBL-EBI
 Protein Domains InterPro IPR008262 Lipase, active site
EMBL-EBI

Trans Membrane

ID	Number Of Domains	Probability of Interior N-Terminus
NP_000220	2	0.05945

Sequence

Target Sequence
 >HUGENEFL:M12625_AT
 cttcaactacacaggccgtgacttccaacgcttctttgcagacctgcactttgaggaagg
 ctggtacatgtggctgcagtcacgtgacctcctggcaggactcccagcacctggtgtgga
 agtatactgtctttacggcgtgggctgcccacgccccgcacctacatctacgaccacgg
 ctccccctacacggacctgtgggtgtgctctatgaggatggtgatgacacggtggcgac
 ccgcagcaccgagctctgtggcctgtggcaggccgccagccacagcctgtgcacctgct
 gcccctgcacgggatacagcatctcaacatggtcttcagcaacctgacctgtgagcacat
 caatgccatcctgctgggtgcctaccgccagggtccccctgcatccccgactgccagccc
 agagccccgcctcctgaataaagaccttctcttctgctaccgtaagccctgatggctatgt
 ttcaggttgaaggaggcactagagtcccacactagggtttcactcctcaccagccacagg
 ctcaagtgtgtgtgcagtg

Probe Sequence(5'-3')	Probe X	Probe Y	Probe Interrogation Position	Strandedness
CTTCAACTACACAGGCCGTGACTTC	152	127	1161	Antisense
CTACACAGGCCGTGACTTCCAACGC	153	127	1167	Antisense
CCAACGCTTCTTTGCAGACCTGCAC	154	127	1185	Antisense
CCTGCACTTTGAGGAAGGCTGGTAC	155	127	1203	Antisense
CATGTGGCTGCAGTCACGTGACCTC	156	127	1227	Antisense
GCTGCAGTCACGTGACCTCCTGGCA	157	127	1233	Antisense
CCTGGCAGGACTCCCAGCACCTGGT	158	127	1251	Antisense
GGACCCTGTGGGTGTGCTCTATGAG	159	127	1353	Antisense
TGTGCTCTATGAGGATGGTGATGAC	160	127	1365	Antisense
GGCGACCCGCAGCACCGAGCTCTGT	161	127	1395	Antisense
CCTGACCCTGGAGCACATCAATGCC	162	127	1503	Antisense
GCACATCAATGCCATCCTGCTGGGT	163	127	1515	Antisense
CATCCTGCTGGGTGCCTACCGCCAG	164	127	1527	Antisense
CTTTGCTACCGTAAGCCCTGATGGC	165	127	1611	Antisense
TACCGTAAGCCCTGATGGCTATGTT	166	127	1617	Antisense
AAGCCCTGATGGCTATGTTTCAGGT	167	127	1623	Antisense
CTATGTTTCAGGTTGAAGGGAGGCA	168	127	1635	Antisense
GGAGGCACTAGAGTCCCACACTAGG	169	127	1653	Antisense
GTCCCACACTAGGTTTCACTCCTCA	170	127	1665	Antisense
CACAGGCTCAGTGCTGTGTGCAGTG	171	127	1695	Antisense

